

Electron paramagnetic resonance studies of Gd³⁺ in K₂Zn(SO₄)₂6H₂O

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Abstract

A single crystal of K₂Zn(SO₄)₂6H₂O doped with Gd³⁺ was studied by EPR at room temperature and at -150°C using an X-band spectrometer. There are two magnetically equivalent spectra characterized by the main Stevens constants $b_2^0 = 0.029 \text{ cm}^{-1}$ and $b_2^2 = 0.0217 \text{ cm}^{-1}$. The spectra are described by a general spin Hamiltonian and all constants have been determined. The zero-field splitting of the energy levels is of the order of 0.3 cm^{-1} and EPR lines exist at very low fields, which are identified. The spectra are temperature dependent. This is interpreted as an anisotropic modification of the Gd³⁺ environment. The analysis of the pseudo-symmetry of the fourth order term of the spin Hamiltonian shows a strong modification of the local structure. The coordination polyhedra of Gd³⁺ are different from Zn and K polyhedra. The comparison with other results from the double sulphate family is made.

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